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What is claimed is:

1. A composition comprising a reduced isoalpha acid (RIAA) and isoalpha acid (IAA) isolated from hops, wherein the RIAA and IAA are in a ratio of about 3:1 to about 1:10.

- 2. The composition of claim 1, wherein said isoalpha acid is selected from isohumulone, isocohumulone, and isoadhumulone.
- 3. The composition of claim 1, wherein said reduced isoalpha acid is selected from dihydro-isohumulone, dihydro-isocohumulone, and dihydro-adhumulone.
- 4. A method of reducing inflammation, comprising administering a composition comprising a reduced isoalpha acid (RIAA) and isoalpha acid (IAA) isolated from hops, wherein the RIAA and IAA are in a ratio of about 3:1 to about 1:10.
- 5. The method of claim 1, wherein said isoalpha acid is selected from isohumulone, isocohumulone, and isoadhumulone.
- 6. The method of claim 1, wherein said reduced isoalpha acid is selected from dihydro-isohumulone, dihydro-isocohumulone, and dihydro-adhumulone.
- 7. A method of reducing inflammation, comprising administering at least two compounds of Genus A having the formula:

wherein R' is selected from the group consisting of carbonyl, hydroxyl, OR, and OCOR, wherein R is alkyl;

and wherein R" is selected from the group consisting of CH(CH₃)₂, CH₂CH(CH₃)₂, and CH(CH₃)CH₂CH₃, wherein the two compounds are in a ratio of about 10:1 to about 1:10.

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AMENDED CLAIMS

received by the International Bureau on 08 August 2005 (08.08.2005): original claims 1-7 have been replaced by amended claims 1-7.

What is claimed is:

- 1. A composition comprising a reduced isoalpha acid (RIAA) and isoalpha acid (IAA) isolated from hops, wherein the RIAA and IAA are in a ratio of about 3:1 to about 1:10 and wherein said RIAA and IAA individually comprise at least 0.1% of the composition.
- 2. The composition of claim 1, wherein said isoalpha acid is selected from isohumulone, isocohumulone, and isoadhumulone.
- 3. The composition of claim 1, wherein said reduced isoalpha acid is selected from dihydro-isohumulone, dihydro-isocohumulone, and dihydro-adhumulone.
- 4. A method for reducing PGE2 mediated inflammation, comprising administering a composition comprising a reduced isoalpha acid (RIAA) and isoalpha acid (IAA) isolated from hops, wherein the RIAA and IAA are in a ratio of about 3:1 to about 1:10 and wherein said RIAA and IAA individually comprise at least 0.1% of the composition.
- 5. The method of claim 4, wherein said isoalpha acid is selected from isohumulone, isocohumulone, and isoadhumulone.
- 6. The method of claim 4, wherein said reduced isoalpha acid is selected from dihydro-isohumulone, dihydro-isocohumulone, and dihydro-adhumulone.

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7. A method FOR reducing PGE2 mediated inflammation, comprising administering at least two compounds of Genus A having the formula:

wherein R' is selected from the group consisting of carbonyl, hydroxyl, OR, and OCOR, wherein R is alkyl;

and wherein R" is selected from the group consisting of CH(CH₃)₂, CH₂CH(CH₃)₂, and CH(CH₃)CH₂CH₃, wherein the two compounds are in a ratio of about 10:1 to about 1:10 and wherein said RIAA and IAA individually comprise at least 0.1% of the composition.